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APPLICATION NO.	FILING DATE	FIRST NAMED INVENTOR	ATTORNEY DOCKET NO.	CONFIRMATION NO.
09/856,415	07/02/2001	James D. Talton	5853-186US	7896
7590	03/30/2004		EXAMINER	
Akerman Senterfitt & Eidon Gregory A Nelson 222 Lakeview Avenue P O Box 3188 West Palm Beach, FL 33402-3188			SHEIKH, HUMERA N	
			ART UNIT	PAPER NUMBER
			1615	
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Please find below and/or attached an Office communication concerning this application or proceeding.

Office Action Summary	Application No.	Applicant(s)
	09/856,415	TALTON ET AL.
Examiner	Art Unit	
Humera N. Sheikh	1615	

-- The MAILING DATE of this communication appears on the cover sheet with the correspondence address --

A SHORTENED STATUTORY PERIOD FOR REPLY IS SET TO EXPIRE 3 MONTH(S) FROM THE MAILING DATE OF THIS COMMUNICATION.

- Extensions of time may be available under the provisions of 37 CFR 1.136(a). In no event, however, may a reply be timely filed after SIX (6) MONTHS from the mailing date of this communication.
- If the period for reply specified above is less than thirty (30) days, a reply within the statutory minimum of thirty (30) days will be considered timely.
- If NO period for reply is specified above, the maximum statutory period will apply and will expire SIX (6) MONTHS from the mailing date of this communication.
- Failure to reply within the set or extended period for reply will, by statute, cause the application to become ABANDONED (35 U.S.C. § 133). Any reply received by the Office later than three months after the mailing date of this communication, even if timely filed, may reduce any earned patent term adjustment. See 37 CFR 1.704(b).

Status

1) Responsive to communication(s) filed on 07 April 2003.

2a) This action is **FINAL**. 2b) This action is non-final.

3) Since this application is in condition for allowance except for formal matters, prosecution as to the merits is closed in accordance with the practice under *Ex parte Quayle*, 1935 C.D. 11, 453 O.G. 213.

Disposition of Claims

4) Claim(s) 28-45 and 48-70 is/are pending in the application.
4a) Of the above claim(s) _____ is/are withdrawn from consideration.

5) Claim(s) _____ is/are allowed.

6) Claim(s) 28-45 and 48-70 is/are rejected.

7) Claim(s) _____ is/are objected to.

8) Claim(s) _____ are subject to restriction and/or election requirement.

Application Papers

9) The specification is objected to by the Examiner.

10) The drawing(s) filed on _____ is/are: a) accepted or b) objected to by the Examiner.

 Applicant may not request that any objection to the drawing(s) be held in abeyance. See 37 CFR 1.85(a).

 Replacement drawing sheet(s) including the correction is required if the drawing(s) is objected to. See 37 CFR 1.121(d).

11) The oath or declaration is objected to by the Examiner. Note the attached Office Action or form PTO-152.

Priority under 35 U.S.C. § 119

12) Acknowledgment is made of a claim for foreign priority under 35 U.S.C. § 119(a)-(d) or (f).
a) All b) Some * c) None of:
1. Certified copies of the priority documents have been received.
2. Certified copies of the priority documents have been received in Application No. _____.
3. Copies of the certified copies of the priority documents have been received in this National Stage application from the International Bureau (PCT Rule 17.2(a)).

* See the attached detailed Office action for a list of the certified copies not received.

Attachment(s)

1) Notice of References Cited (PTO-892)
2) Notice of Draftsperson's Patent Drawing Review (PTO-948)
3) Information Disclosure Statement(s) (PTO-1449 or PTO/SB/08)
Paper No(s)/Mail Date _____.
4) Interview Summary (PTO-413)
Paper No(s)/Mail Date _____.
5) Notice of Informal Patent Application (PTO-152)
6) Other: _____.

DETAILED ACTION

Status of the Application

Receipt of the Change of Address filed 11/19/02, the Supplemental Information Disclosure Statement (IDS) filed 12/03/02, the Request for Extension of Time (3 months-granted) filed 04/07/03, the Amendment filed 04/07/03 and the Interview Summary dated 08/01/03 is acknowledged.

Applicant's arguments as stated in the Interview Summary filed 08/01/03, with respect to the Restriction requirement after a Non-Final Office Action, were fully considered and were found to be persuasive. The Restriction requirement of 07/14/03 has been *withdrawn*.

The Rejections and Election of Species requirement made in the Office Action filed 10/03/02 have been *withdrawn* in view of the 'state of the art'. Accordingly, new ground(s) of rejection follows.

Claims 28-45 and 48-70 are pending. Claims 28, 29, 31, 50, 66-68 and 70 have been amended. Claims 46-47 have been cancelled as requested. Claims 28-45 and 48-70 are rejected.

Note: Applicant's amendment of 04/07/03 includes cancellation of claims 45-47. The Examiner believes that only claims 46-47 were intended to be cancelled based on non-elected species by Applicant in the response filed 04/07/03 and this may be an inadvertent typographical error, since claim 45 recites a "continuous" coating layer and

not "discontinuous". Page 4, lines 17-18 of the response filed 04/07/03, indicates the cancellation of only claim 46 and its dependent, claim 47. Clarification is requested.

Double Patenting

The nonstatutory double patenting rejection is based on a judicially created doctrine grounded in public policy (a policy reflected in the statute) so as to prevent the unjustified or improper timewise extension of the "right to exclude" granted by a patent and to prevent possible harassment by multiple assignees. See *In re Goodman*, 11 F.3d 1046, 29 USPQ2d 2010 (Fed. Cir. 1993); *In re Longi*, 759 F.2d 887, 225 USPQ 645 (Fed. Cir. 1985); *In re Van Ornum*, 686 F.2d 937, 214 USPQ 761 (CCPA 1982); *In re Vogel*, 422 F.2d 438, 164 USPQ 619 (CCPA 1970); and, *In re Thorington*, 418 F.2d 528, 163 USPQ 644 (CCPA 1969).

A timely filed terminal disclaimer in compliance with 37 CFR 1.321(c) may be used to overcome an actual or provisional rejection based on a nonstatutory double patenting ground provided the conflicting application or patent is shown to be commonly owned with this application. See 37 CFR 1.130(b).

Effective January 1, 1994, a registered attorney or agent of record may sign a terminal disclaimer. A terminal disclaimer signed by the assignee must fully comply with 37 CFR 3.73(b).

Claims 68-70 are rejected under the judicially created doctrine of obviousness-type double patenting as being unpatentable over claims 1-36 of U.S. Patent No. 6,406,745 B1 (Talton) in view of Lowndes *et al.* (US Pat. No. 5,499,599). Although the conflicting claims are not identical, they are not patentably distinct from each other because similar subject matter has been claimed. Instant claim 68 is drawn to a method of preparing a medicament, by providing a plurality of core drug particles, having an average particle size of less than 500 μm in diameter, and depositing onto the surface of plurality of core drug particles, at least a first coating layer that comprises a plurality of polymeric coating particles, said coating layer being biodegradable,

biocompatible, wherein the average thickness of the coating layer is between 1 and 500 nm, said depositing step by a process comprising pulsed laser ablation under vacuum, wherein said vacuum is between 1 mTorr and 1 Torr. U.S. Pat. 6,406,745 B1 is also drawn to a method of coating a particulate core material. The only significant distinction observed between instant claims 68-70 and Pat. '745 is that '745 recites a pressure of 'about 10 Torr or higher' whereas instant amended claim 68 recites 'between 1 mTorr and 1 Torr.' The secondary reference of Lowndes *et al.* U.S. Pat. '599 is relied upon for its teaching of a 'method for continuous control of composition and doping of pulsed laser deposition films by pressure control' whereby Lowndes *et al.* teach that the 'Thickness of each layer is controlled by the number of laser shots' (col. 5, lines 27-33) and also teach that 'by controlling the pressure of the gas within the chamber, the composition of the deposit grown upon the substrate can be accurately and continually controlled. Furthermore, by altering gas pressure during the film growth process, the composition of adjacent layers of the film being formed are altered accordingly' (see col. 3, lines 7-12). Therefore, since Patent '599 demonstrates the co-relationship between varying pressure and varying coating thickness, it is deemed obvious to one of ordinary skill in the art to adjust the pressure or Torr in order to obtain the desired or intended coating thickness. Hence, claims 68-70 remain rejected under Non-statutory Obviousness-Type Double Patenting.

Claim Rejections - 35 USC § 103

The following is a quotation of 35 U.S.C. 103(a) which forms the basis for all obviousness rejections set forth in this Office action:

(a) A patent may not be obtained though the invention is not identically disclosed or described as set forth in section 102 of this title, if the differences between the subject matter sought to be patented and the prior art are such that the subject matter as a whole would have been obvious at the time the invention was made to a person having ordinary skill in the art to which said subject matter pertains. Patentability shall not be negatived by the manner in which the invention was made.

The factual inquiries set forth in *Graham v. John Deere Co.*, 383 U.S. 1, 148 USPQ 459 (1966), that are applied for establishing a background for determining obviousness under 35 U.S.C. 103(a) are summarized as follows:

1. Determining the scope and contents of the prior art.
2. Ascertaining the differences between the prior art and the claims at issue.
3. Resolving the level of ordinary skill in the pertinent art.
4. Considering objective evidence present in the application indicating obviousness or nonobviousness.

Claims 28, 30-45, 48, 50-54 and 59-61 are rejected under 35 U.S.C. 103(a) as being unpatentable over Moro et al. (U.S. Pat. No. 5,223,244) in view of Green et al. (US Pat. No. 5,976,577).

Moro et al. teach aerosol compositions comprising at least one propellant, a solvent and a composite powder, whereby a sheath powder, having a particle size of 1/5 or less of a core powder is attached to the core powder that has an average particle size of 0.1 to 100 μm to form a composite powder (see reference column 2, lines 12-44); (col. 4, lines 26-37). The amount of the composite powder is preferably 0.1% to 30% by weight in the total amount of the aerosol composition (col. 5, lines 46-53). In

Example 10, at column 14, lines 6-24, Moro et al. demonstrate the teaching of a powder spray, which comprises an aerosol spray that contains an active ingredient potassium glycyrrhizinate. The composite powder is granular tetrafluoroethylene (1 μm) with a kaolin coating thickness of 0.1 μm . After components (1) to (5) were mixed, the mixture was filled in an aerosol can, followed by filling components (6) and (7) to obtain a powder spray. The spray was found to have a good powder dispersibility and usability.

According to Moro et al., as the core powder of the composite powder usable, is any desired organic powder with a density of 0.7 to 2.0 and an average particle size of 0.1 to 100 μm can be used, and the powder used for the core can be in the form of a spheroid, plate, granule or needle (col. 2, lines 12-19). Moro et al. teach at column 4, lines 35-37, teach that the core powder is substantially completely covered by the coating powder and with a superior stability against separation. As the method of preparing the composite powder, the composite powder can be prepared by mixing the core powder and the sheath powder by the dry process or the wet process (col. 3, lines 33-37).

Moro et al. are deficient in the sense that they do not explicitly teach that the coating layer is a continuous and non-porous layer. It is the Examiner's position that there is no criticality observed in the use of Applicant's continuous or non-porous layer since the instant specification permits the use of both a continuous and discontinuous coating layer (see specification pgs. 2 and 5, lines 29-32). Furthermore, it is deemed obvious to one of ordinary skill in the art to employ a continuous and non-porous coating

layer to obtain a sustained or controlled rate of delivery of the active ingredient. Such skill is also evident from the reference of Green *et al.* (see below).

Green et al. teach a process for preparing rapidly disintegrating solid dosage forms containing drug particles, wherein the drug particles may be *coated or uncoated* with a water-insoluble polymer or lipid material, resulting in a dosage form that exhibits delayed release of the drug for a sufficient period of time to provide for controlled or *sustained release* of the drug after swallowing. The drug particles have a size such that the coatings can be formed thereon which are sufficiently *intact and continuous* to prevent or minimize loss of drug during processing. The coarse drug particles have an average particle size up to about 500 μm . In this size range, it is possible to apply a *uniform intact coating* on the particle in order to achieve efficient freeze-dried dosage forms with slow drug release rate (see reference cols. 1, line 5 – col. 3, line 22); (col. 5, lines 1-47).

Therefore, it would have been obvious to one of ordinary skill in the art at the time the invention was made to use the combined teachings of Moro *et al.* and Green *et al.* because Moro *et al.* teach an active ingredient formulation (i.e., glycyrrhizinate) whereby a core powder is coated and covered by a sheath powder and similarly Green *et al.* teach a dosage form comprising drug particles that are either coated or uncoated and if coated, provide a continuous intact coating on the drug particles in order to provide a sustained release of the active ingredient. The expected result would be a

continuously coated, non-porous pharmaceutical formulation that exhibits a sustained release of the active drug material, as similarly desired by the Applicant(s).

Claims 28, 30-45 and 48-61 and 66-67 are rejected under 35 U.S.C. 103(a) as being unpatentable over Sakon et al. (US Pat. No. 5,972,388) in view of Green et al. (US Pat. No. 5,976,577).

Sakon et al. teach an ultrafine particle powder for inhalation and method for the production whereby the particle powder is produced by spray-drying a mixture of active agent and a lower alkyl ether cellulose wherein the active ingredient and cellulose are either dissolved or suspended in a solution and then spray-dried into particles, whereby 80% of the particles have a particle size in the range of 0.5 to 10 μm . Particles smaller than this size do not appear to be critical since size criticality appears to depend upon administration to lower airways, which is achieved with the teachings of Sakon. Such is also the case for thickness of the coating layer. According to Sakon et al., it is desirable that the medicament is not readily removed by cilia and retained at the site to be deposited. Sustained release of the medicament while it is retained further enhances its efficacy (col. 2, lines 35-40). Sakon et al. teach that the active ingredient includes steroids, such as triamcinolone acetonide and flunisolide, antiallergics, chemotherapy medicaments, antitussives and bronchodilators. These medicaments may be used singly, or as a mixture of two or more thereof unless the mixture is incompatible (col. 9, lines 10-26).

Sakon *et al.* do not teach that the coating layer is a continuous and non-porous layer. It is the Examiner's position that there is no criticality observed in the use of Applicant's continuous or non-porous layer since the instant specification permits the use of both a continuous and discontinuous coating layer (see specification pgs. 2 and 5, lines 29-32). Furthermore, it is deemed obvious to one of ordinary skill in the art to employ a continuous and non-porous coating layer to obtain a sustained or controlled rate of delivery of the active ingredient. Such skill is also evident from the reference of Green *et al.* (see below).

Green et al. teach a process for preparing rapidly disintegrating solid dosage forms containing drug particles, wherein the drug particles may be *coated or uncoated* with a water-insoluble polymer or lipid material, resulting in a dosage form that exhibits delayed release of the drug for a sufficient period of time to provide for controlled or *sustained release* of the drug after swallowing. The drug particles have a size such that the coatings can be formed thereon which are sufficiently *intact and continuous* to prevent or minimize loss of drug during processing. The coarse drug particles have an average particle size up to about 500 μm . In this size range, it is possible to apply a *uniform intact coating* on the particle in order to achieve efficient freeze-dried dosage forms with slow drug release rate (see reference cols. 1, line 5 – col. 3, line 22); (col. 5, lines 1-47).

Therefore, it would have been obvious to one of ordinary skill in the art at the time the invention was made to use the combined teachings of Sakon *et al.* and Green

et al. because Sakon *et al.* teach an ultrafine particle powder formulation comprising medicaments (i.e., triamcinolone acetonide) whereby the particle powder is produced by spray-drying a mixture of active agent and a lower alkyl ether cellulose to provide a sustained release of the medicament and similarly Green *et al.* teach a dosage form comprising drug particles that are either coated or uncoated and if coated, provide a continuous intact coating on the drug particles in order to provide a sustained release of the active ingredient. The expected result would be a continuously coated, non-porous pharmaceutical formulation that exhibits a sustained release of the active drug material, as similarly desired by the Applicant(s).

Claims 28, 30-45, 48, 59-61 and 66-67 are rejected under 35 U.S.C. 103(a) as being unpatentable over Hanes *et al.* (US Pat. No. 5,855,913) in view of Green *et al.* (US Pat. No. 5,976,577).

Hanes *et al.* teach biodegradable aerodynamically light particles incorporating a surfactant on the surface for pulmonary drug delivery whereby the particles are produced by emulsifying active agent in a polymer, such as poly(lactic acid) or PLA; or poly(glycolic acid) or PGA, in a volatile solvent. After mixing, the mixture is spray-dried and the volatile solvent is evaporated to leave the drug particle enclosed within the polymer. The particles are taught to be as small as 2 μm and can also have a mean diameter of between 5 μm and 30 μm . Particles smaller than 2 μm do not appear to be critical since size criticality appears to depend upon administration to lower airways, which is achieved with Hanes *et al.* Such is also the case for the thickness of the

coating layer (see reference col. 5, line 16 – col. 8, line 56) and Abstract. Hanes et al. teach that the aerodynamically light particles are highly suitable for inhalation therapies, particularly in controlled release applications (col. 8, lines 54-56). Various therapeutic agents may be employed in the formulation, including antibiotics and anti-asthmatic agents (col. 10, lines 3-49).

Hanes et al. do not explicitly teach that the coating layer is a continuous and non-porous layer. It is the Examiner's position that there is no criticality observed in the use of Applicant's continuous or non-porous layer since the instant specification permits the use of both a continuous and discontinuous coating layer (see specification pgs. 2 and 5, lines 29-32). Furthermore, it is deemed obvious to one of ordinary skill in the art to employ a continuous and non-porous coating layer to obtain a sustained or controlled rate of delivery of the active ingredient. Such skill is also evident from the reference of Green et al. (see below).

Green et al. teach a process for preparing rapidly disintegrating solid dosage forms containing drug particles, wherein the drug particles may be *coated or uncoated* with a water-insoluble polymer or lipid material, resulting in a dosage form that exhibits delayed release of the drug for a sufficient period of time to provide for controlled or *sustained release* of the drug after swallowing. The drug particles have a size such that the coatings can be formed thereon which are sufficiently *intact and continuous* to prevent or minimize loss of drug during processing. The coarse drug particles have an average particle size up to about 500 μm . In this size range, it is possible to apply a

uniform intact coating on the particle in order to achieve efficient freeze-dried dosage forms with slow drug release rate (see reference cols. 1, line 5 – col. 3, line 22); (col. 5, lines 1-47).

Therefore, it would have been obvious to one of ordinary skill in the art at the time the invention was made to use the combined teachings of Hanes *et al.* and Green *et al.* because Hanes *et al.* teach aerodynamically light drug particles contained within biodegradable polymers that provide for controlled release of the active ingredient for use in pulmonary drug delivery and similarly Green *et al.* teach a dosage form comprising drug particles that are either coated or uncoated and if coated, provide a continuous intact coating on the drug particles in order to provide a sustained release of the active ingredient. The expected result would be a continuously coated, non-porous pharmaceutical formulation that exhibits a sustained release of the active drug material, as similarly desired by the Applicant(s).

Claims 62-65 are rejected under 35 U.S.C. 103(a) as being unpatentable over Moro *et al.* (US Pat. No. 5,223,244) or Sakon *et al.* (US Pat. No. 5,972,388) or Hanes *et al.* (US Pat. No. 5,855,913) in view of Bucks *et al.* (US Pat. No. 6,277,364).

The teachings of Moro *et al.* ('244), Sakon *et al.* ('388) and Hanes *et al.* ('913) have been discussed above. Moro *et al.*, Sakon *et al.* and Hanes *et al.* do not teach the inclusion of a kit having instructions.

Bucks *et al.* ('364) teach aerosol formulations that include kits and packages that comprise labeling instructions for application of the composition for the protection of

skin. The labeling instructions include directions on the amount and frequency of application, methods of removal, suggested storage conditions, shelf life expectancy, precautions or contraindications, and so forth (see reference column 5, line 65 – col. 6, line 7); (claim 4).

Therefore, it would have been obvious for one of ordinary skill in the art at the time the invention was made to use the combined teachings of either Moro et al., Sakon et al. or Hanes et al. with Bucks et al. because Moro et al., Sakon et al. and Hanes et al. all teach powder formulations in aerosol formulations and similarly Bucks et al. teach aerosol formulations that also include kits and packages comprising specific instructions (i.e., method of use, storage conditions, shelf-life extent) for the medicament. The expected result would be effective kit or packaged aerosol formulations that provide for ease and safety of usability.

Claims 29 and 68-70 are rejected under 35 U.S.C. 103(a) as being unpatentable over Moro et al. (US Pat. No. 5,223,244) or Sakon et al. (US Pat. No. 5,972,388) or Hanes et al. (US Pat. No. 5,855,913) in view of Lowndes et al. (US Pat. No. 5,499,599) and further in view of Green et al. (US Pat. No. 5,976,577).

The teachings of Moro et al. ('244), Sakon et al. ('388) and Hanes et al. ('913) are delineated above.

They are lacking in that they do not teach a process of *pulsed laser ablation* and do not explicitly teach a continuous, non-porous coating layer.

Lowndes et al. teach a method for continuous control of composition and doping of *pulsed laser deposited films* by pressure control, wherein by controlling the pressure of the gas within the chamber, the composition of the deposit grown upon the substrate can be accurately and continually controlled. According to Lowndes et al., the thickness of each layer was controlled by the number of laser shots. By varying the gas pressure between two limiting values, and maintaining each pressure for a fixed number of laser shots, structures with highly reproducible layer thickness and composition were fabricated (see column 4, lines 7-18); (col. 5, lines 27-34). Lowndes et al. teach in Fig. 1 at column 2, line 59 – col. 3, line 5, that an ultra-low pressure of about *10 Torr or less* was obtained.

Therefore, it would have been obvious to one of ordinary skill in this art at the time of the invention to use the teachings of Lowndes et al. within Moro et al., Sakon et al. or Hanes et al. because Lowndes et al. explicitly teach that films and layers with varied coating thickness are produced by controlling by the number of laser shots and varying gas pressure using the pulsed laser ablation method and vividly teaches the co-relationship between adjusting pressures and coating thickness and similarly Moro et al., Sakon et al. and Hanes et al. teach formulations comprising an array of coating layers and thicknesses. The expected result would be effective thin film materials obtained by pulsed laser techniques, as similarly desired by the Applicant(s).

As stated previously, Moro *et al.* ('244), Sakon *et al.* ('388) and Hanes *et al.* ('913) do not explicitly teach a continuous, non-porous coating layer.

Green et al. ('577) is relied upon for its teaching of drug particles that may be *coated or uncoated* with a water-insoluble polymer or lipid material, resulting in a dosage form that exhibits delayed release of the drug for a sufficient period of time to provide for controlled or *sustained release* of the drug after swallowing. The drug particles have a size such that the coatings are sufficiently *intact and continuous* to prevent or minimize loss of drug during processing.

Hence, it would have been obvious to one skilled in the art at the time the invention was made to incorporate continuous, non-porous coatings on drug particles if sustained release properties were desired, or discontinuous, porous coatings if rapid release was intended. The expected result would be a particulate/active ingredient formulation with either sustained or immediate release rate properties.

Response to Arguments

Applicant's arguments with respect to claims 28-45 and 48-70 have been considered but are moot in view of the new ground(s) of rejection.

Correspondence

Any inquiry concerning this communication or earlier communications from the examiner should be directed to Humera N. Sheikh whose telephone number is (571) 272-0604. The examiner can normally be reached on Monday through Friday from 7:00A.M. to 4:30P.M.

If attempts to reach the examiner by telephone are unsuccessful, the examiner's supervisor, Thurman Page, can be reached on (571) 272-0602. The fax phone number for the organization where this application or proceeding is assigned is (703) 872-9306.

Any inquiry of a general nature or relating to the status of this application or proceeding should be directed to the receptionist whose telephone number is (703) 308-1235.

hns *TKS*

March 23, 2004

THURMAN K. PAGE
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